

Measuring low activities...

27871
Z/038/61/000/011/002/004
D291/D305

3 tables, 1 figure and 6 references: 5 Soviet-bloc, 1 unidentified.

ASSOCIATION: Ústav jaderného výzkumu, Praha (Nuclear Research Institute, Prague)

Card 4/4

RALKOVA, J.

CZECHOSLOVAKIA

SAIL, J.; RALKOVA, J.

**Nuclear Research Institute, Czechoslovak Academy of Sciences (Institut
für Kernforschung, Tschechoslowakische Akademie der Wissenschaften),
Ros near Prague (for both)**

**Prague, Collection of Czechoslovak Chemical Communications, No 2, Feb
1966, pp 871-880**

**"Behavior of strontium and cesium during the fixation of radioactive
products in basalt."**

RALKOVA, Jarmila

Determination of low specific activities in solutions.
Jaderna energie 8 no.8:281-284 Ag '62.

1. Ustav jaderného výzkumu, Československá akademie věd.

L 48290-65 ENT(m) Feb DIAAP

ACCESSION NR: AP5013585

CZ/0038/65/000/003/0091/0094

AUTHOR: Ralkova, Jarmila

TITLE: Determination of Sr^{90} in water

SOURCE: Jaderna energie, no. 3, 1965, 91-94

TOPIC TAGS: strontium; isotope, water sanitation

ABSTRACT: A rapid control method of Sr^{90} determination that is appropriate if mainly rapidity and less accuracy are required is described. The more accurate methods for the determination of Sr^{90} traces using natural or artificial sorbents are proved on the basis of others' experiences. Orig. art. has: 2 figures, 1 graph.

ASSOCIATION: Ustav jaderného výzkumu CSAV, Rez (Institute of Nuclear Physics, CSAV)

SUBMITTED: 00

ENCL: 00

SUB CODE: NP, GC

NO REF SOV: 000

OTHER: 011

NA

Card 1/1

RALKOVA, Jarmila

Diffusion of radioisotopes in basalt. Silikaty 9 no.1:12-24 '65.

1. Institute of Nuclear Research of the Czechoslovak Academy of Sciences, Rez near Prague. Submitted May 10, 1964.

L 35431-65 EWT(m)/EWP(b)/EWP(e) Pq-4/Peb DIAAP WH

ACCESSION NR: AP5007769

Z/0012/65/000/001/0012/0024

AUTHOR: Ralkova, J.

TITLE: Radioisotope diffusion in basalt ¹⁹ ₁₅

26
21
B

SOURCE: Silikaty, no. 1, 1965, 12-24

TOPIC TAGS: radioisotope disposal, isotope diffusion, basalt, reactor fuel, radioactive waste, glass, aluminum, radiocesium, radiostrontium, isotope fixation

ABSTRACT: Glass and molten rock have been found to be excellent and economical materials to render insoluble highly radioactive wastes, such as spent fuel from reactors which radiates tens of kilocuries/kg and is usually highly acid. Four types of basalt found in Czechoslovakia at Hollberg, Tesetice, Dolanky and Oharice were studied in detail and that from Tesetice was found to absorb ions most rapidly in the resolidified form. The Statni vyzkumny ustav sklarsky (State glass research institute) prepared specimens by melting crushed basalt at 1350-1450C, tempering for two hours at 650C and then cooling gradually for 24 hours to room temperature. Small plates 20 x 20 x 1 mm were cut and ground smooth. Drops of a Cs-137 or Sr-90 solution were placed on such plates and the escape of radioisotope prevented by heating to 600C, cooling, and washing with

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L 35431-65

ACCESSION NR: AP5007769

nitric acid in distilled water. The plates with radioactive cesium were then placed in a small muffle furnace and the heat steadily raised from 440C while measuring the radiation at specific intervals with a beta GM tube. Radiation from similar strontium-90 specimens was measured and recorded after deducting impulses caused by the accompanying Y-90. Each specimen was tested without a filter and then with an aluminum filter of 124 mg/cm², which absorbed all Sr-90 rays and half the Y-90 rays. Radiation rates of the two isotopes in basalt were charted at temperatures from 440 to 850C and found to be on the order of 10⁻³ up to 600C for cesium and 660C for strontium, with 10⁻² at higher temperatures. The diffusion coefficients of each isotope in basalt and in glass were tabulated and charted, as well as their absorption curves in aluminum. As the temperature rose from 440 to 850C, the diffusion coefficients varied from around 10⁻¹³ to 10⁻¹¹. The physical and chemical properties of Tesetice basalt were also studied in relation to heat treatment ranging from 650 to 1000C, and x-ray studies were made of the internal structure after eight types of tempering. Crystal size and the recrystallization process were analyzed in relation to ion diffusion at various temperatures, and the ion activation energy was found to be $E_{Cs} = 95$ kcal/mol and $E_{Sr} = 105$ kcal/mol, which would indicate predominantly interstitial diffusion. The chief advantage of basalt is its recrystallization at high temperatures, which improves its physical and chemical properties for isotope

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L 35431-65

ACCESSION NR: AP5007769

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fixation. If the decay heat does not exceed 600C, Cs and Sr diffusion coefficients will not exceed 10-15 cm²/sec, which is suitable for safe disposal of radioactive wastes. "The author is indebted to Dr. J. Voldan of the Stat. vyzkumny ustav sklarsky (State glass research institute) for providing the basalt specimens and analyzing their properties, Dr. E. Kucera, at the Ustav fysikalni chemie CSAV (Physical chemistry institute) for the x-ray analyses, and V. Vesely for help in preparing the specimens." Orig. art. has: 5 figures, 5 tables and 12 formulas.

ASSOCIATION: Ustav jaderneho vyzkumu CSAV, Rez near Prague (Nuclear research institute, CSAV)

SUBMITTED: 10May64

ENCL: 00

SUB CODE: MT, NP

NO REF SOV: 001

OTHER: 007

Card 3/3

BALOV, R.

Animals of Tanganyika. IUn.nat. no.8:20-21 Ag '57. (MLRA 10:8)
(Tanganyika Preserve)

11 AND 120 CODES										11 AND 120 CODES									
PROCESSES AND PROPERTIES INDEX																			
CA																			
An apparatus for determining warfare agents and in- jurious gases in the air. M. M. Baker, <i>Polish Patent</i> undisclosed (Germany 11, No. 10, 37 0(1940); <i>Chem.</i> <i>Zentr.</i> 1941, 8, 619.—An easily transportable app. is de- scribed. W. R. Henn																			
ASO-5LA METALLURGICAL LITERATURE CLASSIFICATION																			
FROM SYNTHESE										FROM SYNTHESE									
COLLECTOR										COLLECTOR									

54. Sodium hexametaphosphate from technical phosphoric acid, by M. Ralsceva and I. Ránya. ["Magyar Kémiai Folyóirat" Hungarian Journal of Chemistry Vol. 56, No. 5, pp. 201-203, May 1950.]

Technical phosphoric acid, which does not contain free sulfuric acid and possibly little monocalciumphosphate, was carefully neutralized to pH 4.2 with sodium carbonate. The greater portion of the calcium, iron and aluminum impurities precipitated during this treatment. Vanadium did not precipitate in laboratory experiments. In large scale operations, however, the precipitate also absorbed vanadium. Monosodiumphosphate was not crystallized from the obtained filtered solutions, but gradually evaporated to dryness and brought to red heat. This method has the advantage that the impurities can be precipitated during evaporation and be removed by a second filtration at about a 50 Bar degree concentration. The evaporation was performed in a lead container, the glowing in an ordinary fire clay crucible. The obtained sodium hexametaphosphate was white, or slightly greenish when traces of vanadium were present. It showed all the characteristic reactions.

RALTEANU, G.; NASTASEANU, S.

Upper Paleozoic horizons of the Ciudanovita-Lupac (Banat) region.
p. 135.

ANALELE SERIA STINTELOR NATURII. Bucuresti, Rumania. Vol. 7, no. 18, 1958.

Monthly List of East European Accessions (EEAI), LC, Vol. 8, no. 9, Sept., 1959.

Uncl.

PALETTE, J.

[illegible]

RALIS, Zdenek

SURNAME, Given Names

Country: Czechoslovakia

Academic Degrees: MD

Affiliation: Clinic of Surgery, Faculty of Medical Hygiene KU /Karlova univer-
sita; Charles University/ (Chirurgicka klinika lekarske fakulty
hygienicke KU), Prague 10; Director: Prof E. POLAK, MD.

Source: Prague, Prakticky Lekar, Vol 41, No 9, 1961, pp 387-391.

Data: "Relapsing Nephrolitiasis."

Authors: VACH, Bohumil

RALIS, Zdenek

260
GPO 981643

SIRBILADZE, N.Ya.; RALISHVILI, L.T.; DROZDOVA, Ye.; MYL'NIKOVA, T.A.; KARCHKHADZE, R.G.

Production of pyrogen-free antidiphtheria and antitetanus therapeutic sera. Nauch. osn. proizv. bakt. prep. 10:196-205 '61. (MIRA 18:7)

1. Tbilisskiy institut vaktsin i syvorotek.

RALJ-100, 10

Ralićević, S. Sur certaines classes de polynômes et sur la répartition de leurs zéros. Srpska Akad. Nauka. Zb. Rad. 50, Mat. Inst. 5 (1956), 1-60. (Serbo-Croatian. French summary)

L'objet de ce travail est l'interprétation géométrique des transformations linéaires (affines) effectuées dans les polynômes de la forme

$$(*) \quad \sum_{k=0}^v \frac{n}{n-k} \binom{n-k}{k} (-p^2)^k z^{n-2k} + q, \quad v = [n/2],$$

et en particulier la répartition des zéros des polynômes transformés ainsi obtenus. Par des considérations géométriques élémentaires, l'auteur établit que la construction des zéros des polynômes transformés se déduit avec la même précision que celle des polynômes réguliers correspondants aux (*). La démonstration d'une suite de résultats de ce genre repose sur la notion et les propriétés de "l'écart" d'un polynôme $P_n(z)$ défini par

$$R = \max |P_n(\zeta_v)| - \min |P_n(\zeta_v)|, \quad P_n'(\zeta_v) = 0$$

$$(v = 1, 2, \dots, n-1).$$

M. Tomić (Beograd).

On a Certain Class of Polynomials and on the Distribution of their Zeros.

RALIK, Josef, MUDr; URBANOVA, Slavomira, MUDr

Glycaemic curve in patients with keratoconjunctivitis sicca. Cesk.
ofth. 10 no.5:321-325 Oct 54.

1. Z I. oční kliniky Karlovy university v Praze. Prednosta doc.
Dr. E.Dienstbier

(KERATOCONJUNCTIVITIS

sicca, glucose in)

(GLUCOSE TOLERANCE TEST, in various diseases
keratoconjunctivitis sicca)

RALIN, E. M.
N. V. BFLOV, Russ. 35,320, Mar. 31, 1934

RALITSKAYA, S. I.

"The Clinical Treatment of Atypical Mastoiditis." Cand Med Sci, First Moscow
Order of Lenin Medical Inst, 1 Nov 54. (VM, 20 Oct 54)

Survey of Scientific and Technical Dissertations Defended at USSR Higher
Educational Institutions (10)

SO: Sum. No. 481, 5 May 55

RALIZIN, G. S.

G. S. Ralizin, "On Some Applications of Matrix Algebra to the Theory of Mechanisms."

paper presented at the 2nd All-Union Conf. on Fundamental Problems in the Theory of Machines and Mechanisms, Moscow, USSR, 24-28 March 1978.

RAIJAVIO, A.

Correction to the work "A Straight and a Characteristic Segment in Polygons
with Zeros of Polynomials." p. 114.
ZHURNAL MATEMAT. GEOMETRII, No. 43, 1955.

SD: Monthly List of East European Accessions, (SOL), 1955, Vol. 1, No. 10, Oct. 1955,
Encl.

✓ Raljević, S. Sur certaines classes de polynômes et sur la répartition de leurs zéros. Srpska Akad. Nauka. Zb. Rad. 50, Mat. Inst. 5 (1956), 1-60. (Serbo-Croatian. French summary)

L'objet de ce travail est l'interprétation géométrique des transformations linéaires (affines) effectuées dans les polynômes de la forme

$$(*) \quad \sum_{k=0}^v \frac{n}{n-k} \binom{n-k}{k} (-p^2)^k z^{n-2k} + q, \quad v = [n/2],$$

et en particulier la répartition des zéros des polynômes transformés ainsi obtenus. Par des considérations géométriques élémentaires, l'auteur établit que la construction des zéros des polynômes transformés se déduit avec la même précision que celle des polynômes réguliers correspondants aux (*). La démonstration d'une suite de résultats de ce genre repose sur la notion et les propriétés de "l'écart" d'un polynôme $P_n(x)$ défini par

$$R = \max |P_n(z_v)| - \min |P_n(z_v)|, \quad P_n'(z_v) = 0 \quad (v=1, 2, \dots, n-1).$$

M. Tomić (Beograd).

On a Certain Class of Polynomials and on the Distribution of their Zeros.

2

Raljević, S. Remarque sur un théorème de M. Marden.
Srpska Akad. Nauka. Zb. Rad. 55, Mat. Inst. 6 (1957), 16
69-72. (Serbo-Croatian. French summary)

This note shows by example that Theorem (44.1) is incorrect as stated in the reviewer's "The geometry of the zeros of a polynomial in a complex variable", Math. Surveys, no. 3, Amer. Math. Soc., N.Y., 1949 [MR 11, 101], p. 157. As the reviewer has known for some time, the statement in the theorem " $f(z)$ has $n-k$ zeros on the unit circle" should read " $f(z)$ has $n-k$ zeros on or symmetric in the unit circle". The other zeros mentioned in the theorem are in addition to these zeros. See also F. F. Bonsall and M. Marden, Proc. Amer. Math. Soc. 3 (1952), 471-475; 5 (1954), 111-114 [MR 13, 938; 15, 613].

M. Marden (Milwaukee, Wis.)

5/1

RALJEVIC, S.

SCIENCE

Periodical: ZBORNIK RADOVA. No. 55, 1957.

RALJEVIC, S. A remark on a Marden's theorem. p. 69.

Monthly List of East European Accessions (EEAL) LC, Vol. 2, no. 3
March 1957 Unclass.

1973, 1.

"A Generalization of Euler's Vector and ..."
(SIAM J. MATH. ANAL. 11, v. 1, no. 1, 1973,
Zagreb, Yugoslavia)

SO: Monthly Bulletin of East European Academics, Moscow, October, 1973,
no. 10, October, 1973, Unclassified

RAIKO, A.V.

Kinetics of firing processes. Inzh.-fiz.zhur. no.5:12-16 My '60.
(MIRA 13:8)

1. Ordena V.I.Lenina politekhnicheskij institut, Kiyev.
(Ceramics)

Ralko, A.V.

USSR/Atomic and Molecular Physics - Heat

D-4

Abs Jour : Ref Zhur - Fizika, No 1, 1958, 768

Author : Ralko, A.V.

Inst : -

Title : Analysis of Investigations of Non-Stationary Heat Exchange and Mass Exchange in Phase Transformations and in Chemical Transformations.

Orig Pub : Tr. mosk. tekhnol. in-t pishch. prom- sti, 1957, vyp. 8, 215-219

Abstract : No abstract.

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S/170/60/003/005/002/017
B012/B056

/c

AUTHOR: Ralko, A. V.

TITLE: The Kinetics of Baking Processes

PERIODICAL: Inzhenerno-fizicheskiy zhurnal, 1960, Vol. 3, No. 5,
pp. 12 - 16

TEXT: With a view to working out methods for the purpose of automatically obtaining optimum curves for clay-baking, the author experimentally investigated the kinetics of clay-baking. In a paper (Ref. 2), the author described a method in which three differential thermocouples were distributed over the experimental sample. This made it possible, by mathematical evaluation of the experimental thermograms obtained, to determine the quantitative thermophysical characteristics and the thermal effects of the investigated mineral (Ref. 3). Fig. 1 shows three thermograms which were obtained when baking Prosyanyaya kaolin. The thermograms were recorded according to the above method by an electronic multipoint potentiometer. These experimental data show that the chemical transformations in the heating of a body first take place in its surface

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The Kinetics of Baking Processes

S/170/60/003/005/002/017
B012/B056

layer (zone of chemical reactions). Later, the zone of chemical transformations gradually shifts into the depth of the body. The calculations dealt with here confirm the mechanism of the gradual deepening of the zone of chemical reactions in the capillary-porous body. The zone of chemical reactions gradually shifts into the interior according to the parabolic law, which agrees with the Stefan theorem on the "freezing-through" of moist bodies. The rapid increase of the temperature gradient during the shifting of the zone of the endothermic effect causes thermal stresses in the sample, which may lead to the destruction of the material. An artificial slowing down of the development of the endothermic effect at a given temperature gradient on the boundary of chemical transformations in workpieces can be attained only by a corresponding decrease of the heat supply from outside during baking. Fig. 2 shows the circuit diagram of the experimental arrangement which is described. The experiments are briefly described, and Fig. 3 shows the curves obtained in the baking of clay plates having a thickness of 20 mm when heated to a temperature of 900°C. These curves distinctly show the endothermic effect in kaolin. By means of the method shown here for the determination of baking-curves, optimum modes of operation in the baking of workpieces

Card 2/3

The Kinetics of Baking Processes

S/170/60/003/005/002/017
B012/B056

from various ceramic materials may be determined. The nature of the method consists in the fact that the electronic controller automatically brings about the necessary furnace conditions by means of a prescribed mode of operation necessary for the respective sample and material. This means that it forms the optimum curve for baking. There are 3 figures and 5 Soviet references. ✓c

ASSOCIATION: Ordena V. I. Lenina politekhnicheskii institut, g. Kiyev
("Order of V. I. Lenin" Polytechnic Institute, Kiev)

Card 3/3

RALKO, A. V.

" Study of heat - and mass- exchange in a process of roasting
clays and kaolins."

Report presented at the 1st All-Union Conference On Heat- and Mass Exchange,
Minsk, BSSR, 5-9 June 1961

RALKO, A. V.

Experimental Investigation of Unsteady Heat- and Mass-Transfer During
Phase and Chemical Transformations

Akademiya nauk SSSR. Energeticheskiy institut
Teplo- i massoobmen v protsessakh ispareniya (Heat- and Mass-Transfer in
Evaporation Processes) Moscow, Izd-vo AN SSSR, 1958. 254p. 5,000 copies
printed.

RAL'KO, A. V.; KURSENKO, I.

"Thermodynamics of irreversible heat and mass transfer as a scientific method for studying the kinetics of the firing of silicates and silicate articles."

report submitted for 2nd All-Union Conf on Heat & Mass Transfer, Minsk, 4-12 May 1964.

Kiev Polytechnic Inst

Fuller
RALKO, A.V.

Analysis of studies on unstable heat and mass transfer in phase
and chemical transformations. Trudy MTIPP no.8:215-219 '57.

(MIRA 10:12)

(Mass transfer) (Heat--Transmission)

RAIKO, A. V.

RAIKO, A. V. --"Experimental Investigation of Nonstationary Heat and Mass Exchange during Phase and Chemical Conversion." Min of Higher Education USSR, Moscow Technological Inst of Food Industry, Moscow, 1956 (Dissertation for the degree of Doctor of Technical Sciences.)

KNIZHNYI LETOPIS
No 41, October 1956

RALKO, A. V.

"Investigation of Heat and Mass Transfer at the Process of Baking
of Clays and Caolins."

Report submitted for the Conference on Heat and Mass Transfer, Minsk,
BSSR, June 1961.

RAL'KO, V.A., Geroy Sotsialisticheskogo Truda; LOBANOV, A.P.; KURLYPO, M.F.;
YANUSHEVSKAYA, M.S.; FEDOTKINA, A.I.

Introducing scientific farm management on the "Stalin" Collective
Farm. Zemledelie 7 no.8:6-11 Ag '59. (MIRA 12:10)

1.Predsedatel' kolkhoza imeni Stalina, Pinskogo rayona, Brestskoy
oblasti (for Ral'ko). 2.Nauchno-issledovatel'skiy institut ekono-
miki i organizatsii sel'skokhozyaystvennogo proizvodstva Akademii
sel'skokhozyaystvennykh nauk BSSR (for Lobanov, Kurlypo, Yanu-
shevskaya). 3.Belorusskiy nauchno-issledovatel'skiy institut zhi-
votnovodstva (for Fedotkina).

(White Russia--Collective farms)

CHUMAKOV, Yu.I.; LEDOVSKIKH, V.M.; LOKHOV, R.Ye.; BALKO, V.A.

1,3-di-(2-pyridyl)propane. Metod.poluch.khim.reak. i prepar. no.7:
56-57 '63. (MIRA 17:4)

1. Kiyevskiy politekhnicheskii institut.

BAKANOV, A.; RAL'KOV, V.

Small-size moistening apparatus with automatic feeding of water.
Muk.-elev. prom. 29 no.12:20-21 D '63. (MIRA 17:3)

1. Nachal'nik razmol'nogo tsekha Tul'skogo mel'nichnogo kombinata No.1 (for Bakanov).
2. Zamestitel' nachal'nika razmol'nogo tsekha Tul'skogo mel'nichnogo kombinata No.1 (for Ral'kov).

RUL'KO, V.I., Gercy Priblislavitskogo Truda

land has become more generous. Philadelphia 27 no. 01 78-78 N '01.
(MIRA 18'10)

1. Predsedatel' kollektiva 'Denezhitskiy', Pinskogo rayona, Krestakoy
oblasti.

RALKOVA, Jarmila

Diffusion of radioisotopes in glass. Silikaty 6 no.3:258-272 '62.

1. Ustav jaderného výzkumu, Řez u Prahy.

RALKOVA, Jarmila

Measurement of radioactivity in water. Jaderna energie 6 no.3:89-93
Mr '60.

1. Ustav jaderného výzkumu, Praha.

L 33603-65 EWT(m)/EPF(c)/EPF(n)-2/ENG(m)/EPR Pr-4/Ps-4/Pu-4

ACCESSION NR: AP5009492

Z/0038/65/000/001/0009/0011 **32**

AUTHOR: Berak, Lubomir; Dlouhy, Zdenek (Dlougi, Z.); Kepak, Frantisek; Napravnik, Jiri (Napravnik, Y.); Ralkova, Jarmila (Ralkova, Ya.); Saidl, Jaroslav (Saidl, Ya.); Schejbalova, Ludmila (Sheybalova, L.); Vesely, Vladimir (Vesel', V.); Zaruba, Josef (Zaruba, Y.)

TITLE: Problems of radioactive wastes being solved in the Institute of Nuclear Research of the Czechoslovak Academy of Sciences

SOURCE: Jaderna energie, no. 1, 1965, 9-11

TOPIC TAGS: radioactive waste disposal, radioactive waste disposal equipment

ABSTRACT: Investigations of radioactive waste disposal are reviewed. Some methods developed and proved are briefly described. Several of them became a basis for pilot plant and full operation equipment design and construction. Orig. art. has: 3 figures.

ASSOCIATION: Ustav jaderneho vyzkumu CSAV, Rez (Institute for Nuclear Research, CSAV)

SUBMITTED: 00

ENCL: 00

SUB CODE: NP

NO REF SOV: 000

OTHER: 017

NA

Card 1/1

BALEK, Jaroslav, Ing. Dr. Sc., BALKOVA, Jarmila, Dr. Sc.; BLUNDOKO, Jaroslav,
Ing.

Use of Isotopes for Hydrologic Measurements. Vodni hosp 15 no.4:
145-147, 1969.

1. Institute of Hydrodynamics of the Czechoslovak Academy of
Sciences, Prague, and Institute of Nuclear Research of the
Czechoslovak Academy of Sciences, 1962.

22363
Z/038/61/000/001/004/005
A201/A126

215300

AUTHORS: Rálková, Jarmila and Slunečko, Jaroslav

TITLE: Measurement of low activities in waters. III. Design of a cell-type counter

PERIODICAL: Jaderná energie, no. 1, 1961, 13 - 15

TEXT: The article describes the improvements made in a cell-type scintillation counter described by the same authors in a previous article. The counter was developed at the Ústav jaderného výzkumu ČSAV (Institute of Nuclear Research, ČSAV) and has been used for some time for the measurement of waste-water activity at the nuclear reactor in Řež. The first major improvement concerns the cell design. The original flat cell, which was made entirely of scintillation material with a volume of 2 milliliters, was replaced by a larger cell with a volume of 20 milliliters. Its walls are of polished plexiglas and inside the cell, thin, polyvinyl-toluene-base plastic scintillator plates, produced by Tesla Liberec, are mounted at 1 mm intervals parallel to the axis of two photomultipliers, as shown in Figure 1. This arrangement has the following advantages: The self-absorption in the sample is reduced; the geometric efficiency is as high as 94%; the back-

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Z/038/61/000/001/004/005
A201/A126

Measurement of low activities ...

ground is reduced to 0.8 count/min due to the reduced total mass of the scintillation material; and larger samples can be measured. The new design of the cell has improved the measuring efficiency by a 1.5 order of magnitude so that activities as low as 5×10^{-11} c/liter can be measured without previous thickening. With tenfold thickening, water samples of a specific activity of 5×10^{-12} c/liter can be measured, which is in compliance with the Czechoslovak standard ČSN 567900 for drinking water. The cell is inserted into two plexiglas lightpipes connected optically with two US-made RCA 6342 photomultipliers. The wiring diagram of the high-voltage divider and cathode follower is shown in Figure 2. Positive high voltage is tapped separately for each photomultiplier from two Tesla scintillation detectors. The photocathode is grounded. Scintillation pulses are taken from the last dynode. Output pulses are fed through a shielded cable to amplifiers which are built-in in the Tesla detectors. The three-stage amplifiers have an input sensitivity of 20, 80, or 300 mv. Output pulses are positive with a height of 25-30 v. These shaped pulses are fed into the coincidence circuit. The coincidence circuit has a resolution time of 10^{-7} sec and was designed by the elektronické oddělení ÚJV (Electronics Section, ÚJV). It eliminates all noise pulses generated in the photomultipliers and lets pass through only pulses which were detected by both photomultipliers simultaneously. The pulses from the coincidence circuit are

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A201/A126

Measurement of low activities ...

positive, have a height of 10 v and a duration of 0.5 microseconds. They are then fed to the anticoincidence circuit. The anticoincidence shielding of the scintillation counter is formed by a double ring of STS 8 GM tubes, and its purpose is to reduce the background due to cosmic rays. If a cosmic particle still penetrates through the GM-tube ring and reaches the scintillator, the produced scintillation pulses are detected by both photomultipliers simultaneously so that they pass through the coincidence circuit and have to be separated from pulses produced by the radiation of the sample. This is done by a special anticoincidence circuit the wiring diagram of which is shown in Figure 3, and which constitutes the second major improvement. The anticoincidence function proper is performed by the gating circuit E3E4. Before entering the gating circuit, pulses are shaped by shaping circuits. Output pulses from the coincidence circuit are fed over input S0 and trigger the monostable flip-flop circuit E1E2 formed by the double triode ECC85. The generated, negative, square pulses with a height of about 40 v and a duration of 25μ sec are derived by the CR circuit so that two short pulses, one negative and one positive, are obtained. The positive pulse is delayed by about 25μ sec and has a duration of 1.5μ sec. The GM-pulses are negative, of longer duration and a slow build-up time. They are fed over the input GM to the amplifying stage E6 with a reduced anode voltage and short transfer characteristic.

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Z/038/61/000/001/004/005
A201/A126

Measurement of low activities ...

Thus, positive pulses with a steep leading edge are obtained in the anode circuit of the triode. These pulses are then further derived by the derivating circuit CR and their positive peaks trigger the monostable flip-flop circuit E7E8 formed by the double triode ECC85, while the negative peaks are clipped by the germanium diode 4NN41. From the flip-flop circuit, rectangular pulses with a height of about 15 v and a duration of 35 μ sec are fed to the grid of the triode E4 of the gating circuit. The gating circuit E3E4 is formed by the double triode ECC85. Both triodes have a common resistance of 4 k Ω . The triode E4 is opened by grid bias tapped from a 160 k Ω and a 6.4 k Ω voltage divider. A current flows through this vacuum tube and the cathode resistance causing a voltage drop of about 30 v so that the triode E3 is closed. Shaped pulses from the input Sc are fed to the control grid of the triode E3. Only a delayed positive pulse with a height of 35 v opens this vacuum tube. The amplified voltage pulse is tapped from the 20 k Ω anode resistance and power-amplified by the cathode follower E5. The output pulses are negative, with a height of 40 v and a duration of 1.5 μ sec. When a shaped GM pulse is fed to the grid of the triode E4 of the gating circuit E3E4, the voltage rises on the cathode resistance of 4 k Ω from 30 to about 45 v, since the triode E4 is connected as cathode follower. The triode E3 remains closed so that none of these pulses can appear on the output circuit of the instrument. When a pulse from

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Z/038/61/000/001/004/005
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Measurement of low activities ...

the input Sc (40 v, 1.5 μ sec, delayed by 25 μ sec) appears on the grid of the triode E3 at a time when on the grid of the triode E4 there is still a GM pulse (15 v, 35 μ sec), this pulse is stopped by the gating circuit since its height is not sufficient to open the triode E3 at which the bias has just been increased to 45 v. Thus it is ensured that pulses which were produced simultaneously in GM tubes and the scintillator are separated from pulses produced in the scintillator alone. The instrument in this arrangement has proved to be successful in operating conditions. Appreciation is extended to Václav Veselý for drawing the diagrams and assisting in building the instrument. There are 3 figures and 2 Soviet-bloc references.

ASSOCIATION: Ústav jaderného výzkumu ČSAV (Institute of Nuclear Research),
Prague

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84639

Z/038/60/000/007/003/006
A201/A026

26.2246
AUTHORS:

Rádková, Jarmila; Slunéčko, Jaroslav

TITLE:

Low-Activity Measurements in Waters. II. The Design of a Cell Counter

PERIODICAL: Jaderná energie, 1960, No. 7, pp. 235 - 237

TEXT: The Ústav jaderného výzkumu ČSAV (Institute of Nuclear Research, CSAV) developed a cell-type scintillation counter for measuring low-level β - activity in liquids. It consists of a flat cell, made of hydrophobic plastic scintillator, mounted between two photomultiplier tubes in coincidence connection, as shown in Figure 1. This assembly is surrounded by a ring of STS 8 G M tubes, as shown in Figure 2, which are connected in anticoincidence relative to the scintillation counter. This anticoincidence envelope absorbs cosmic radiation thus reducing the natural background of the counter. The entire assembly is enclosed in a 10-cm lead shield to keep off external γ -radiation. High-tension voltage for the photomultipliers is supplied from a source similar to that used in Tesla scintillation counters, with a built-in three-stage amplifier of input sensitivities 0.02, 0.08 and 0.3 v. Since the pulses from the photomultipliers have a height of about 100 mv, an input sensitivity of 0.08 v is quite adequate.

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Low-Activity Measurements in Waters. II. The Design of a Cell Counter

Output pulses have heights between 25 and 30 v, the amplifier stability being 0.1% at $\pm 10\%$ grid-voltage changes. The coincidence circuit was designed by the elektronické oddělení ÚJV (Electronics Section, ÚJV). Power for GM tubes and cathode followers is supplied from a stabilized Tesla BS 275 power source. The decade scaler was produced by the Leninovy závody (Lenin Works) in Plzeň. US RCA 6342 photomultipliers were used, but Czech or Soviet photomultipliers can be used, too. A special anticoincidence circuit was designed for this counter by the Institute of Nuclear Research. Its main part is a gating circuit which has to meet the following requirements: 1) To let pass all scintillation pulses, which passed the coincidence circuit and did not originate simultaneously with any of the GM tube pulses. 2) To arrest those scintillation pulses which originated simultaneously with some of the GM tube pulses. 3) To arrest all pulses which originated in the GM tubes. Prior to entering the gating circuit, the GM tube pulses are shaped to a rectangular pattern and the scintillation pulses are delayed in such a manner that both pulse types, if they originated simultaneously, meet at the gating-circuit input. This prevents pulses, caused in the scintillator by a cosmic particle, from appearing at the output of the apparatus.

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Low-Activity Measurements in Waters. II. The Design of a Cell Counter

The block-schematic of the apparatus is shown in Figure 3. The apparatus has a background of 2 - 3 pulses/min (with a 10-cm lead shield). Its geometry-efficiency is 80% with a cell having inside dimensions of 20 x 30 x 3 mm. The activity of several samples of the Vltava River water, the potable water from the municipal water system and reactor water was measured and the results compared to data obtained by GM counters, as shown in Table 1. It was found that the cell-type scintillation counter makes possible direct measurements of liquids with a specific activity $\sim 10^6 \mu\text{c/ml}$, with an error of $\pm 12\%$. The liquid measured must not attack the scintillation material and has to be transparent to blue and green light. When larger quantities of sample liquids are condensed, even liquids with lower activities can be measured, down to $10^{-9} \mu\text{c/ml}$ (reactor water) with a maximum error of $\pm 30\%$. Measuring time is 1 hour for samples yielding less than 5 pulses/min, and 30 minutes for samples yielding higher counts. Background is measured during 1 hour prior to and after sample measuring. The measuring efficiency of the counter can be further increased by modification of the cell. Experiments are being conducted with a cell composed of several plastic scintillator plates spaced 1 mm apart parallel to the axis of the photomultipliers. The

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Low-Activity Measurements in Waters. II. The Design of a Cell Counter

geometry-efficiency of this cell is 94%. Scintillation pulses are led to photo-cathodes so that even opaque liquids can be measured. Preliminary calculations indicate that activity of river and potable water as low as $10^{-9} \mu \text{ c/ml}$ can be detected. Credit is given to Doctor J. Juna for his suggestions and V. Veselý and M. Petrň for their help in building the counter and sample preparation. There are 2 photographs, 1 figure, 1 table, and 11 references: 2 Czech, 8 English and 1 German. (Edited by I. Bučina and J. Juna).

ASSOCIATION: Ústav jaderného výzkumu ČSAV, Praha (Institute of Nuclear Research, ČSAV, Frague)

Card 4/4

Diffusion of

Z/012/62/000/003/001/001
E073/E335

500	1.7×10^{-11}
550	2.5×10^{-11}
600	2.9×10^{-11}

The values for strontium in the temperature range 350 - 575 °C were:

350	3.07×10^{-13}
400	5.10×10^{-13}
450	1.95×10^{-12}
500	1.26×10^{-11}
525	1.73×10^{-11}
550	1.89×10^{-11}
575	2.20×10^{-11}

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Diffusion of

Z/012/62/000/003/001/001
E073/E335

The activation energy of diffusion changes with temperature and this is in accordance with the hypothesis on the activation of the glass structure. The diffusion coefficients of caesium are lower than those of other alkalis with smaller ionic diameters. However, the activation energy of diffusion of Cs is lower than that of sodium and this is considered proof of the assumption that caesium is the most weakly bound element in glass. Caesium migrates faster than strontium at lower temperatures, due to the fact that in this range fusion is affected by the valency of the ions. In the region of softening of the glass the diffusion coefficients of mono- and bivalent ions become approximately equal. The low values of the diffusion coefficients indicate that the sealing of highly radioactive elements into glass would be safe provided the rate of elution were limited solely by diffusion. Results of earlier elution tests with fused basalt indicate that in the initial stage the surface layers are flushed and eluted and even dissolved at a rate higher by several orders of magnitude than that corresponding to diffusion. Later on, the elution factors are of an

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Diffusion of

Z/012/62/000/003/001/001
E073/E335

order of magnitude comparable with that of the diffusion coefficients. Further work is scheduled on the diffusion of ions in types of fused basalt which are considered to be chemically most stable and economically most favourable. There are 8 figures and 7 tables.

ASSOCIATION: Ústav jaderného výzkumu, Řež u Prahy
(Institute of Nuclear Research, Řež, Prague)

SUBMITTED: August 28, 1961

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RALKOVA, Jarmila; SLUNECKO, Jaroslav

Low activity measurement in waters. Part 2: The design of a cell counter. Jaderna energie 6 no.7:235-238 J1 '60.

1. Ustav jaderného výzkumu, Československá akademie věd, Praha

RALKOVA, Jarmila

Low activity measurement in water; part 4. Jaderna energie 7 no.11:
374-377 N '61.

1. Ustav jaderného výzkumu, Praha.

VALEVA, Jarmila; VALL, Jaroslav

Incorporation of radioisotopes into melted silicates.
Jaderna energie 10 no.1114-18 Ja'64.

1. Ústav jaderného výzkumu, Československá akademie věd, Pz.

31923
Z/038/62/000/002/004/004
D286/D303

21.5140

AUTHOR:

Rádková, Jarmila

TITLE:

Radioisotope diffusion in glass

PERIODICAL:

Jaderná energie, no. 2, 1962, 56-57

TEXT:

This is an abstract of the 17-page Report No. 540 (12 May 61) of the ÚJV (Nuclear Research Institute). The report, intended as contribution to solving problems encountered in radio-active waste disposal, investigates the diffusion of Cs^{134} and Sr^{90} in glass at 500 - 600°C. The radioisotopes were spread on thin glass plates, the plates were heated, and beta energies were measured on both sides of the plate. The calculated diffusion coefficients ranged from $D = 10^{-13}$ to 10^{-11} cm^2/sec^{-1} . The changes of the diffusion activation energy in dependence of time are in agreement with the hypothesis of glass-lattice activation. The assumption that Cs is the alkali metal with the weakest bond to glass could be confirmed. It was also found that diffusion coefficients of Cs and Sr

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Radioisotope diffusion in glass

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differ only slightly at lower temperatures (glass softening), but greatly at higher temperatures (575°C) which, in turn, confirms the Blau theory stating that fields of force, i.e. ion valences, are the primary factor in diffusion processes. It is finally stated that measured diffusion coefficients permit the assumption that highly-active waste may safely be fixed in glass. There are 7 figures in the original report.

Card 2/2

X

RÁLKOVÁ, JARMILO

CM The polarographic reduction of formaldehyde in the mixture ethanol-water. Jarmila Rálková (Charles Univ., Prague). *Chem. Zvesti* 9, 401-40 (1965). The kinetic wave of reduction of CHO in buffered aq. solns. of EtOH was studied. The presence of EtOH decreased considerably the height of the limiting current which could be explained by the decrease of flow of free reducible aldehydic form of EtOH to the electrode. Jan Micko

ACC NR: AR6034095

SOURCE CODE: UR/0089/66/021/004/0285/0289

AUTHOR: Saidl, Y^{aroslav}; Ralkova, Yarmila (A)

ORG: Institute of Nuclear Research, Czechoslovak Academy of Sciences, Řež near Prague, Czechoslovakia (Institut jaderných issledovaní Chekchoslovatskoy akademii nauk, Řež near Prague, Czechoslovakia)

TITLE: Solidification of radioactive waste by fusing into basalt

SOURCE: Atomnaya energiya, v. 21, no. 4, 1966, 285-289

TOPIC TAGS: radioactive waste disposal, silicon compound, mineral, radiation shielding, physical diffusion

ABSTRACT: With an aim at finding means of isolating highly radioactive products from the biosphere, the authors first review earlier similar work done in Czechoslovakia on the incorporation of various mixtures of fission products in materials with a silicon base. Basalt was chosen because of its chemical stability and mechanical strength, hydrolytic stability and high melting temperature, as well as great abundance in Czechoslovakia. Brief reports are presented on the effect of impurities on basalt, of tests of its radiation endurance, and on the volatility of radioactive isotopes from it. Tests have shown that recrystallization of the vitreous phase of fused basalt improves its properties, primarily its mechanical and chemical endurance. At temperatures 30 - 70C, the calculated diffusion coefficients range from 10^{-15} - 10^{-17} cm²/sec. It is concluded that fused basalt satisfies all requirements imposed on materials used for safe and economic isolation of radioactive waste with large

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UDC: 621.039.7

ACC NR: AP6034095

specific activity. Orig. art. has: 1 figure and 1 table.

SUB CODE: 18/ SUBM DATE: 25Apr66/ ORIG REF: 006/ OTH REF: 011

Card 2/2

ACC NR:AP6021984

(A,N)

SOURCE CODE: UR/0375/66/000/004/0020/0026

AUTHOR: Pirumov, V. S. (Captain 2d Rank); Rall', D. S. (Engineer; Captain 1st Rank; Candidate of Naval Sciences); Trukhayev, R. I. (Engineer)

ORG: None

TITLE: Decision making theory and the control of forces

SOURCE: Morskoy sbornik, no. 4, 1966, 20-26

TOPIC TAGS: operations research, military personnel, cybernetics, command and control system, electronic computer, *NAVAL FORCE ORGANIZATION*

ABSTRACT: The application of cybernetics to the theory of command decisions from the naval viewpoint is discussed. The decision-making process is broken down into... four stages: (1) the formation of an approximate qualitative model; (2) the formalization of the assigned mission; (3) formal optimization; (4) and creative optimization, the last stage prior to the decision itself. The third stage is the most complex, and involves a quantitative optimization of a formal constituent on the basis of an elucidation of the nature and volume of the parameters characterizing the knowledge of the elements of the situation. Levels of knowledge of situation parameters are distinguished and strategic level is examined in somewhat more detail. An example of the application of the theory is given, and it is

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ACC NR: AF6021984

pointed out that the sequence of the process as shown can serve as the basis for further improvement of the command structure and points the way to a degree of possible interaction between personnel and electronic computers in an automated command system. Orig. art. has: 1 table.

SUB CODE: 05,12/SUBM DATE: None

Card 2/2

RALL', D.S., kand. voyenno-morsk. nauk, dotsent, inzh.-kapitan 1-go ranga,
TRUKHAYEV, R.I.

Dynamic programming and certain fields of its use. Mor. sbor. 48 no.11:
21-29 N '64. (MIRA 18:1)

RALLI, D.S., kand. voyenno-morskikh nauk, inzh.-kapitan 1-go ranga; NIKITIN, I.M.,
inzh.-kapitan 2-go ranga; TRUKHAYEV, R.I.

Using the game method for making an optimal decision under conditions
of uncertainty. Mor. sbor. 47 no.3:27-35 Mr '64. (MIRA 18:7)

RALL, J.M.

Are there intelligent beings in the universe? Elet tud 16 no.27:840-
842 2 J1 '61.

RAIL, K.B.; PEREKALIN, V.V.

Reaction of diketenes with some aromatic hydroxy compounds. Zhur. Priklad.
Khim. 25, 1330-5 '52. (MLRA 5:12)
(CA 47 no.21:11155 '53)

1. A.I. Gertsen State Pedagog. Inst., Leningrad.

RALL', K. B.

"Reaction of Diketene With Some Aromatic Oxy-Compounds."
Cand Chem Sci, Leningrad State Pedagogical Inst, Leningrad,
1954. (RZhKhim, No 21, Nov 54)

Survey of Scientific and Technical Dissertations Defended at USSR
Higher Educational Institutions (11)

SO: Sum. No.521, 2 Jun 55

RALL, K.D.

Reactions of diketene. III. Reaction of diketene with some aromatic hydroxy compounds. Synthesis of phenyl acetates. K. B. Rall and V. V. Perekalin (A. I. Gertsen Pedagogical Institute, Leningrad). *Zhur. Obshch. Khim.* 25, 278-81; *J. Gen. Chem. (U.S.S.R.)* 25, 289-91 (1955) (Engl. translation); cf. *C.A.* 50, 300h. To 4.7 g. PhOH in 30 ml. C_6H_6 was added a little pyridine followed, at -10° , by dropwise addn. of 4.5 g. diketene (I) over 10 min.; after refluxing 1.5 hrs. the mixt. yielded 72.3% $AcCH_2CO_2Ph$ (II), m. 50° ; a 63.9% yield was obtained similarly in Et_2O in the presence of few drops of H_2SO_4 . II (1.78 g.) in 100 ml. $EtOH$ and 10 ml. 0.1N $AcONa$ was treated with cooling over 2 hrs. with 110 ml. 0.1N PhN_2Cl , yielding 61% PhO_2CCH_2Ac (III); yellow, m. $63-6^\circ$; this refluxed with 5% HCl gave 86% $AcCH_2(CO_2H)N:NPh$, m. 155° . II in concd. H_2SO_4 gave after 20 hrs. at room temp. a low yield of 4-methylcoumarin, m. 80° . $o-MeOC_6H_4OH$ (5.2 g.) in C_6H_6 , treated with a little pyridine, followed by 4.5 g. I gave after 1.5 hrs. refluxing 68% $o-MeOC_6H_4O_2CCH_2Ac$, m. $62-3^\circ$ (with H_2SO_4 as catalyst the yield was 58.6%); this with PhN_2Cl , as above, gave 83% $o-MeOC_6H_4O_2CCH_2N:NPh$, yellow, m. $95-7^\circ$. Hydroquinone (3.3 g.) and 2.8 g. I, in C_6H_6 in the presence of pyridine, gave after 2 hrs. at reflux and extn. with hot $CHCl_3$ a 91% yield of $p-HOC_6H_4O_2CCH_2Ac$, m. 102° , which is readily hydrolyzed with hot 5% $NaOH$. The use of an excess of I gave $p-C_6H_4(O_2CCH_2Ac)_2$ (III), m. $121-2^\circ$; the same formed from I and the mono deriv. in 84% yield. III and PhN_2Cl , as above, gave 97% $p-C_6H_4(O_2CCH_2AcN:NPh)_2$, orange, m. $226-7^\circ$ (from Ac_2O). $p-MeC_6H_4OH$ and I in C_6H_6 with a little pyridine similarly gave 92.5% $p-MeC_6H_4O_2CCH_2Ac$, m. 69° . Similarly was obtained 74.5% $p-BrC_6H_4O_2CCH_2Ac$, m. 51° . I (0.84 g.) added over 1.5 hrs. to 1.2 g. salicylaldehyde and 0.1 g. dry $KOAc$ at $35-40^\circ$, then treated with 5% HCl , gave 33.7% 3-acetylcoumarin, m. $120-1^\circ$. G. M. Kosolapoff

MA
MST
①

RALL, K.B.

CH Reactions of diketene. IV. Reaction of diketene with some aromatic hydroxy compounds. Synthesis of methylcoumarin derivatives. K. B. Rall and V. V. Perekalin (Gertsen Pedagog. Inst., Leningrad). *Zh. Org. Khim.* 25, 815-21; *J. Gen. Chem. U.S.S.R.* 25, 781-6(1965) (Engl. translation); cf. *C.A.* 50, 1704c.—In reactions of aromatic HO compds. with diketene (I) basic catalysts aid the formation of acetoacetyl derivs., while acid catalysts aid the formation of coumarins. This is readily explained by different form of polarization of I in the initial stage of reaction. To 5.5 g. resorcinol in 50 ml. Et₂O was added a few drops H₂SO₄ and then over 10 min. at -10° 5 g. I and the mixt. refluxed 2.5 hrs. to give 54.5% 7-hydroxy-4-methylcoumarin, m. 186-7°. Phloroglucinol similarly gave 88.5% 5,7-dihydroxy-4-methylcoumarin, m. 233-7°. Pyrogallol gave 7,5-dihydroxy-4-methylcoumarin, m. 235-6°, while hydroxyhydroquinone gave 6,7-dihydroxy-4-methylcoumarin, m. 268-9°. To 7.2 g. 1-C₁₀H₇OH in 75 ml. C₆H₆

was added few drops pyridine followed by 4.5 g. I; after refluxing 1.5 hrs. the mixt. gave 77.2% 1-naphthyl acetoacetate (II), m. 73°; this heated with 5% NaOH gave Me₂CO and 1-naphthol. II treated with PhN₂Cl in the presence of NaOAc gave 1-C₁₀H₇O₄CCHAcN: NPh, red, m. 112-15°. I, 1-C₁₀H₇OH, and a little H₂SO₄ in Et₂O gave 44.6% 4-methyl-7,8-benzocoumarin, m. 160-7°, the same forming in 40% yield from II on heating with AlCl₃ in PhNO₂ 1 hr. at 60°. After 43 hrs. at room temp., 2-C₁₀H₇OH, I, and a little pyridine in C₆H₆ gave 79.6% 2-C₁₀H₇O₄CCHAc, m. 82-3°; this heated with 5% NaOH gave 3-C₁₀H₇OH; coupling with PhN₂Cl in NaOAc soln. gave 98% orange 2-C₁₀H₇O₄CCHAcN: NPh, m. 132-4°. G. M. K.

(1)

BALL, K. B

5

✓ New method of preparation of coumarin derivatives.
K. B. Ball and V. V. Perekalin (A. I. Gertsch State Pedagog.
Inst., Leningrad). *Doklady Akad. Nauk S.S.S.R.* 100,
1715-17 (1955); cf. C.A. 48, 9813t. To 1.2 g. o-HOC₆H₄-
CHO and 0.1 g. KOAc at 35-40° was added in 1.5 hrs. 0.84
g. diketene, yielding 40.4% 3-acetylcoumarin, m. 121-2°.
Phloroglucinol (0.63 g.) in 50 ml. Et₂O at -10° treated with
a little H₂SO₄ and 1 g. diketene and the mixt. refluxed 1.5 hrs.
gave 88.5% 5,7-dihydroxy-4-methylcoumarin, m. 286-7°
(from MeOH). 1-C₆H₄OH (7.2 g.) in C₆H₆ treated at -10°
with a little pyridine and 4.5 g. diketene and refluxed 1.5
hrs. gave 72.6% 1-C₆H₄-O-CCH₃Ac (I), m. 72°. The same
mixt. (with a little H₂SO₄ instead of pyridine) refluxed 2.5
hrs. gave 44.6% 4-methyl-7,8-benzocoumarin, m. 167-8°,
also formed in 40% yield from 1.14 g. I in PhNO₂ treated
with 0.88 g. AlCl₃, kept 48 hrs., and heated 1 hr. at 60°.
G. M. Kosolapoff

① AA 324

RA LL K B

Conjugated systems. LXII. Condensation of hydrocarbons with propiolic acid and its methyl ester. A. A. Petrov and K. B. Rall (Inst. Aviation Instr., Lenin grad). *Zhur. Obshchei Khim.* 26, 1588-93 (1950); cf. C.A. 50, 3292c, 11930a. — Heating 3 g. $\text{HC}\equiv\text{CCO}_2\text{H}$ with 6 ml. $\text{CH}_2=\text{CHCH}=\text{CH}_2$ in 10 ml. MePh with 0.1 g. hydroquinone in a sealed tube 10 hrs. at $145-50^\circ$ gave 88% 2,5-dihydrobenzoic acid, m. 121° , which failed to condense with maleic anhydride. The acid heated with S 45 min. gave BzOH , while treatment with Br in CHCl_3 gave a brominated acid, m. $151-2^\circ$. Heating 6.5 g. $\text{HC}\equiv\text{CCO}_2\text{Me}$ with 10 ml. $\text{CH}_2=\text{CHCH}=\text{CH}_2$ in MePh with 0.1 g. hydroquinone 6 hrs. at 120° gave 70% Me 2,5-dihydrobenzoate, b. $94-5.5^\circ$, d. 1.3720 , n_D^{20} 1.4938, which sapon. with 10% NaOH to the acid described above. Heating 4.8 g. $\text{HC}\equiv\text{CCO}_2\text{H}$ and 12 g. piperylene in MePh with 0.1 g. hydroquinone 12 hrs. at 120° gave 74% 2-methyl-2,5-dihydrobenzoic acid, m. $85-6^\circ$, which failed to condense with maleic anhydride and heated with S gave BzOH ; heating with 20% HNO_3 to 200° gave phthalic acid. Similar reaction of $\text{CH}_2=\text{CHCH}=\text{CH}_2$ with $\text{HC}\equiv\text{CCO}_2\text{Me}$ gave in 10 hrs. at $140-5^\circ$ 71% Me 2-methyl-2,5-dihydrobenzoate, b. $89-90^\circ$, d. 1.0269 , n_D^{20} 1.4871, which with 10% NaOH gave the free acid, m. $85-6^\circ$, identical with the above. Heating 3.5 g. $\text{HC}\equiv\text{CCO}_2\text{H}$ with 3.5 g. isoprene and 0.1 g. hydroquinone in MePh 10 hrs. at $145-50^\circ$ gave 78% 4-methyl-2,5-dihydrobenzoic acid (I), m. $180-1^\circ$, which failed to condense with maleic anhydride on simple heating, but heated with it in the presence of H_2O and Pd black 6 hrs. it gave $p\text{-MeC}_6\text{H}_4\text{CO}_2\text{H}$; oxidation with HNO_3 gave terephthalic acid. Heating I in MeOH in the presence of H_2SO_4 gave the Me ester, b. 110.5° , d. 1.0430 , n_D^{20} 1.4920; the same ester, b. $110.5-11.5^\circ$, d. 1.0435 , n_D^{20} 1.4930, formed in 73% yield from 6 g. $\text{HC}\equiv\text{CCO}_2\text{Me}$ and 7 ml. iso-

2

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Petrov, A. A. and KALL, K. B.

prene heated in MePh with 0.1 g. hydroquinone 10 hrs. at 140-5°. Attempted hydrogenation of the ester over Pd gave a mixt. of partly and completely aromatized esters; hydrogenation did proceed over Raney Ni. Heating 5 g. HC:CCO₂H with 0 g. cyclohexadiene in MePh in the presence of 0.1 g. hydroquinone 7 hrs. at 115-20° gave some C₂H₄ and 40% BzOH. Heating 3.7 g. HC:CCO₂H with 4.2 g. 2-chlorobutadiene in MePh with 0.1 g. hydroquinone 12 hrs. at 115-20° gave 72% 4-chloro-2,5-dihydrobenzoic acid, m. 210-17°, which with dil.HNO₃ at 155° in 10 hrs. gave p-ClC₆H₄CO₂H. LXIII. Action of N,N-dichlorobenzenesulfonamide on alcoholic solutions of piperylene. T. A. Zyryanova and A. A. Petrov (Lensovet Technol. Inst., Leningrad). *Zhur. Obshchei Khim.* 26, 1500 601(1956); cf. C.A. 50, 3990d. To 135 g. piperylene in 600 ml. MeOH was added in 3-4 hrs. at 0-5° 135 g. PhSO₂NCl₂; after standing until active Cl was nearly absent, the mixt. was steam distd. in the presence of a little Na₂SO₄. Fractionation of the distillate gave unreacted piperylene, 2 g. crude chloropentadiene, b. 100-7°, and 43.8 g. of a mixt. (I) of MeCH:CHCH(OMe)CH₂Cl and MeCH(OMe)CH:CHCH₂Cl, b. 55-60°, d₂₀ 0.9784-08.56, n_D²⁰ 1.4431-1.4451 (the products were partially sepd. by fractionation). Ozonolysis gave AcOH and ClCH₂CO₂H. Refluxing the ethers with KOH in MeOH 18 hrs. gave MeCH:CHCH(OMe)CH₂, b. 105-7°, d₂₀ 0.8350, n_D²⁰ 1.4535 (with 5% H₂SO₄ this gave MeCOCCH:CHMe), and MeCH(OMe)CH:CHCHOMe, b. 62.5-3.5°, d₂₀ 0.8882, n_D²⁰ 1.4220, which hydrogenated over Pd to the *sald. analog*, b. 141.5-5°, d₂₀ 0.8381, n_D²⁰ 1.4050. Hydrogenation of I over Raney Ni in the presence of CaCl₂ in MeOH gave a series of fractions containing: MeCH(OMe)CH:CHCHOMe, b. 88-5°, d₂₀ 0.9011, n_D²⁰ 1.4380, and 100 papers specimen of MeCH(OMe)CH:CHCHOMe which hydrogenated

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over Pd gave the pure *sald. ether*, b. 88-8.5°, d₂₀ 0.7516, n_D²⁰ 1.3840; an authentic specimen was prep'd. from MeI and KOH heated with *sec*-amyl alc. in a sealed tube at 100°; the product, b. 88-8.5°, d₂₀ 0.7504, n_D²⁰ 1.3840. Hydrogenation of I also gave a mixt. (Ia) of MeCH:CHCH(OMe)CH₂Cl and PrCH(OMe)CH₂Cl (II), b₂₀ 48-8.5°, which hydrogenated over Raney Ni at 70 atm. H yielded pure II ether, b. 143-4°, b₂₀ 48-8.5°, d₂₀ 0.9590, n_D²⁰ 1.4250; bromination of Ia gave some 14% pure II, which heated with KOH in Cellosolve 4 hrs. gave a little 2-methoxy-1-pentene (III), b. 89-90°, d₂₀ 0.7888, n_D²⁰ 1.4062; this shaken with 5% H₂SO₄ gave MeCOPr, which confirmed the structure. Ozonolysis of III gave PrCO₂H. Reaction of 68 g. piperylene in 200 ml. EtOH with 87.5 g. PhSO₂NCl₂ as described above gave a mixt. (IV) of MeCH:CHCH(OEt)CH₂Cl (IVa) and MeCH(OEt)CH:CHCH₂Cl, partially sepd. in fractions, b₂₀ 50-81°, d₂₀ 0.9570, n_D²⁰ 1.4390, and b₂₀ 81-4°, d₂₀ 0.9813, n_D²⁰ 1.4427; these were combined and exam'd.; ozonolysis gave AcOH and ClCH₂CO₂H; refluxing with EtOH-KOH 18 hrs. gave 2-ethoxy-1,3-pentadiene, b. 124-6°, d₂₀ 0.8294, n_D²⁰ 1.4500 (with 5% H₂SO₄ this gave MeCOCH:CHMe), and MeCH(OEt)CH:CHCH₂OEt (IVb), b₂₀ 76.5-7°, d₂₀ 0.8564, n_D²⁰ 1.4235. IVb hydrogenated over Pd in MeOH to MeCH(OEt)CH₂CH₂CH₂OEt, b₂₀ 73-6°, d₂₀ 0.8511, n_D²⁰ 1.4102. Hydrogenation of IV over Raney Ni in MeOH at 120 atm. H gave mixed MeCH(OEt)CH:CHMe and MeCH(OEt)Pr (V), b. 106-7° (the mixt. hydrogenated over Pd gave the pure V, b. 105-6°, d₂₀ 0.7536, n_D²⁰ 1.3880, identical with a specimen from *sec*-amyl alc. and EtI with KOH), and a mixt. of 1-chloro-2-ethoxypentane (VI) and IVa, b₂₀ 60-1°; further hydrogenation of the latter mixt. over Ni gave the pure VI, b₂₀ 60-60.5°, d₂₀ 0.9359, n_D²⁰ 1.4230. The results indicate that in the reaction of piperylene with PhSO₂NCl₂ the molar ratio of products of 3,4-addn. and 1,4-addn. is 5-6 to 4. Thus, the presence of a Me group in piperylene greatly increases the yield of 1,4-adducts, with fixation of the Cl atom on the 4-position.

G. M. Kosolapoff

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RALL, K. B.

✓ Conjugated systems. LXII. Condensation of diene by-
drocarbons with propiolic acid and its methyl ester. A.A.
Petrov and K. B. Rall. *J. Gen. Chem. U.S.S.R.* 26, 1779-81
(1950) (English translation).—See C.A. 51, 1887e. LXIII.
Action of *N,N*-dichlorobenzene-sulfonamide on alcoholic
solutions of piperylene. T. A. Zyryanova and A. A.
Petrov. *Ibid.* 1785-03.—See C.A. 51, 1887i. B.M.R.

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PEUROV, A.A.; RALL', K.B.

Conjugated systems. Part 81: Order of adding bromine to
dialkyl-vinylacetylenyl- carbinols. Zhur. ob. khim. 27 no.9:
2402-2406 S '57. (MIRA 11:3)

1. Leningradskiy institut aviatsionnogo priborostroyeniya.
(Bromination) (Alcohols)

RLL 13

79-2-32/64

AUTHORS:

Petrov, A. A., Mall', K. B., Semenov, G. I.

TITLE:

Investigations in the Field of Conjugated Systems
(Issledovaniya v oblasti sopryazhennykh sistem). LXXXIII. The
Structure and Reactivity of the Methyl Ether of Vinylacetylene
Carboxylic Acid (LXXXIII. Stroyeniye i reaktsionnaya sposob-
nost' metilovogo efira vinilatsetilenkarbonovoy kisloty).

PERIODICAL:

Zhurnal Obshchey Khimii, 1958, Vol. 28, Nr 2, pp. 424-428 (USSR)

ABSTRACT:

In the series of investigations of reactions at compounds with
double and triple bindings the reaction of the above mentioned
compound with bromine was investigated. The bromization takes
place very slowly, is, however, accelerated at light (graphic-
ally shown). Since the disintegrates dibromide obtained decays
in vacuum distillation the structure of the compound obtained
was investigated in the infrared spectrum. For this purpose al-
so the infrared spectra of the initial compound and of some
other ethers of similar structure were taken. The methyl ether
was obtained according to the prescription of I. N. Nazarov
and M. V. Kuvartzina, the others according to the usual methods.
Data concluding the spectrometer and the bromization are given.
The measurements obtained showed that the bromization takes
place at the double bindings and the compound thus remains an

Card 1/2

Investigations in the Field of Conjugated Systems
LXXXIII. The Structure and Reactivity of the Methyl Ether of Vinylacetylene
Carboxylic Acid.

7-2-32/64

acetylene compound. Explanations of the various infrared spectra obtained and the connections between the different spectral bands and the structure of the compounds are given. There are 2 figures, 1 table, 10 references, 8 of which are Slavic.

ASSOCIATION: Technological Institute imeni Lensovet, Leningrad (Leningradskiy tekhnologicheskii institut imeni Lensovet)

SUBMITTED: February 20, 1957

AVAILABLE: Library of Congress

Card 2/2

PETROV, A.A.; FILL', K.B.; VIL'DAVSKAYA, A.I.

Preparation of nitrodienes and nitroenynes. Zhur. ob. khim. 34
no.10:3513-3514 0 '64. (MIRA 17:11)

1. Leningradskiy tekhnologicheskij institut imeni Lensoveta.

L 44159-65 EPF(c)/EWP(j)/EWA(c)/EWT(m)/T Pc-4/Pr-4 RM

ACCESSION NR: AP5009018

S/0366/65/001/002/0240/0243

AUTHORS: Petrov, A. A.; Rall', K. B.; Vil'davskaya, A. I.

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β

TITLE: Synthesis and properties of nitroalkadienes-1,3 7

SOURCE: Zhurnal organicheskoy khimii, v. 1, no. 2, 1965, 240-243

TOPIC TAGS: nitroalkadiene, hydrocarbon, nitrogen dioxide, iodine

ABSTRACT: Various nitroalkadienes were produced by dehydrohalogenating the product of combining diene hydrocarbons with nitrogen dioxide and iodine, as described by K. B. Rall' and A. A. Petrov (ZhOKh, 34, 3621 1964). The yield of nitroalkadienes was 10-35%, with the major part of the matter suffering resination. All the nitroalkadienes represented yellow lacrimator liquids distillable in vacuum. Their structures and purity were determined from infrared nuclear magnetic resonance spectra. An attempt to produce 1-nitro-2-chlorbutadiene by separating hydrogen iodide from the product of combining NO₂ and iodine with chloroprene proved unsuccessful. The experimental part of the work is presented in detail, and the nitroalkadienes obtained are listed and described. Orig. art. has: 1 formula, 2 graphs, and 1 table.

Card 1/2

L 44159-65

ACCESSION NR: AP5009018

ASSOCIATION: Leningradskiy tekhnologicheskij institut imeni Lensovata (Leningrad
Technological Institute)

SUBMITTED: 22Jan64

ENCL: 00

SUB CODE: 00

NO REF SOV: 004

OTHER: 002

Card 2/2 *116*

RALL', K.B.; PETROV, A.A.

Reaction of diene hydrocarbons with nitrogen tetroxide and
iodine. Zhur. ob. khim. 34 no.11:3621-3624 N '64 (MIRA 18:1)

1. Leningradskiy tekhnologicheskii institut imeni Lensoveta.

RALL', K.B.; PETROV, A.A.

Conjugated systems. Part 154: Vinylacetylene derivatives of mercury. Zhur.ob.khim. 32 no.4:1095-1097 Ap '62. (MIRA 15:4)

1. Leningradskiy tekhnologicheskii institut imeni L'nsoveta.
(Mercury) (Butenyne)

RALL', M.B.

Practical significance of the combined use of cholecystography and
duodenal catheterization in the diagnosis and therapy of cholecystitis.
Vest. khir. 85 no. 7:109-117 Je '60. (MIRA 14:1)
(GALL BLADDER--RADIOGRAPHY) (DUODENUM--EXAMINATION)

RALL', M.B., Cand Med Sci -- (diss) "Acute cholecystitis
and ~~the~~ remote consequences of its treatment." Len, 1950,
15 pp (First Len Med Inst im akademika I.P. Pavlov. Chair
of General Surgery) 200 copies (KL, 50-6, 130)

- 142 -

EXCERPTA MEDICA Sec 9 Vol 13/11 Surgery Nov 59

6797. REMOTE RESULTS IN THE TREATMENT OF PATIENTS WITH ACUTE CHOLECYSTITIS (Russian text) - Rall M. B. - KHIRURGIYA 1956, 11 (52-58) Tables 5

Of 638 patients with acute cholecystitis, 117 were operated upon, while 521 were treated conservatively. The operative mortality amounted to 7.6%. The results of treatment were studied in 222 patients (60 operated on and 162 treated conservatively). Remote results of surgical treatment with follow-up of from 1 to 10 yr. were good in 38 patients, satisfactory in 16 and poor in 6. The results of the conservative treatment were good in 30, satisfactory in 22 and poor in 110 patients. These data show that surgical treatment of acute cholecystitis gives better remote results than conservative therapy. However, it is not expedient to employ surgical treatment in all cases. Patients in whom the clinical picture is not very pronounced have to be subjected to examination of the bile ducts, after subsidence of the acute symptoms, further treatment depending on the results of examination.

RALL', H.B.

Remote results of treating acute cholecystitis. Khirurgia, Moskva 34
no.11:52-58 N '58. (MIRA 12:1)

1. Iz kafedry obshchey khirurgii (zav. - chlen-korrespondent AMN SSSR
prof. A.W. Filatov) i Leningradskogo meditsinskogo instituta im. I.P.
Pavlova.

(CHOLECYSTITIS, ther.
acute, remote results (Rus))

RALL', M.B.

Analysis of the causes of unfavorable outcomes in the surgical
treatment of cholecystitis. Khirurgiia 36 no.4:63-68 Ap '60.

(MIRA 13:12)

(GALL BLADDER--SURGERY)

RALL', M.B., kand.med.nauk; ZARETSKIY, I.P.

Diagnostic significance of cholecystotomography. Vest. khir. 92 no.3:
56-59 Mr '64. (MIRA 17:12)

1. Iz kliniki obshchey khirurgii (zav. - prof. S.M.Kurbangaleyev) 1-go
Leningradskogo meditsinskogo instituta imeni Pavlova.

RALL', S. F. Dr. Tech. Sci.

Dissertation: "Production of Nutritive Glucose, and Crystallization as a Principal Process of its Obtaining." Inst. of National Economy, imeni G. V. Plekhanov, 24 Oct 47.

SO: Vechernyaya Moskva, Oct, 1947 (Project (#17636)

Rall, Yu. M.

20638 Rall, Yu. M. Otstayushchaya oblast biologicheskogo obrazovaniya. [Zoologiya pozvonochnykh]. Vestnik vyssh. shkoly, 1949, No. 6, s. 36-37

SO: LETNIE ZHURNAL STATY - Vol. 28, Moskva, 1949

Otstayushchaya oblast' biologicheskogo obrazovaniya [Zoologiya pozvonochnykh]
Lagging Regional Biological Education [Zoology of the spine]

Vestnik vyssh. Shkoly (Herald of Higher Schools)

RALL', Y.M.

Lec 52

USSR/Geophysics - Tsimlyansk Project

"In the Sands of Tsimlyanskiy Peninsula," Prof Y.M. Rall'

Priroda, No 12, pp 100-105

States that the creation of the Tsimlyanskoye Sea has formed new peninsulas and bays in the tremendous water reservoir, which has posed new problems in the transformation of nature. Zoologists of Rostov U made their ecological survey of the sandy lands as early as May, Jun 52.

263T90

RAIL', Yu.M.; KRITSKAYA, T.I.

Result of the acclimatization of Ussuri raccoon dogs in Rostov Province.
Zool.zhur. 32 no.3:513-523 My-Je '53. (MLRA 6:6)

1. Rostovskiy gosudarstvennyy universitet imeni V.M. Molotova.
(Rostov Province--Raccoon dogs)

RAIL', Yu.M.

Nature and animals of the Tsimlyansk Peninsula. Izv. Vses. geog. ob-va 85
no. 4:382-392 J1-Ag '53. (MLR 6:8)
(Tsimlyansk Peninsula--Natural history) (Natural history--Tsimlyansk
Peninsula)

RAIL, Yu.M.; KARAYEROVA, G.P.

New data on the occurrence and the harmful activity of the
Jerboa Scirtopoda telus Licht. Zool.shur. 33 no.5:1184-1185
S-O '54. (MLRA 7:11)

1. Rostovskiy gosudarstvennyy universitet im. V.M.Molotova.
(Rostov Province--Jerboas) (Jerboas--Rostov Province)

RAIL', Yu.M.; KIYANOVA, V.S.; STRELINA, T.D.

Observations of rodents in irrigated fields in Rostov Province. Zoel.shur. 33 no. 6:1390-1395 M-D '54. (MIRA 8:2)

1. Restevskiy gosudarstvennyy universitet im. V.M.Molotova.
(Rostov Province--Rodentia)

L
RALL', Yu.M.

Animal herd life as an example of intraspecific relationships. Bot.
zhur.40 no.4:565-567 J1-Ag'55. (MLRA 8:11)

1. Protivozhumnyy institut, g. Stavropol'-Kavkazskiy
(Animals, Habits and behavior of)